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(54) **Combinations of CRF antagonists and renin-angiotensin system inhibitors**

(57) The present invention relates to compositions and methods of achieving a therapeutic effect including, but not limited to, the treatment of congestive heart failure or hypertension in an animal, preferably a mammal including a human subject or a companion animal, using a corticotropin releasing factor (CRF) antagonist and a renin-angiotensin system (RAS) inhibitor.

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Description**Background Of The Invention**

5 [0001] The present invention relates to compositions and methods of achieving a therapeutic effect, preferably including the treatment of congestive heart failure or hypertension in an animal, preferably a mammal including a human subject or a companion animal, using a corticotropin releasing factor (CRF) antagonist, preferably in synergistic combination with a renin-angiotensin system (RAS) inhibitor.

[0002] In preferred embodiments of the compositions and methods of this invention, the CRF antagonists comprise 10 compounds of structural formula I or II, including the pharmaceutically acceptable salts thereof, as defined hereinbelow.

[0003] The compounds of formula I or II, their pharmaceutically acceptable salts, and methods of preparing such compounds and salts are disclosed in commonly assigned PCT publication numbers WO 95/34563, WO 95/33750, WO 94/13676, WO 94/13677 and European patent application number EP 773 023. Each of the applications corresponding to PCT publication numbers WO 95/34563, WO 95/33750, WO 94/13676 and WO 94/13677 designate, inter alia, the 15 United States and are incorporated herein by reference in their entirety.

[0004] The foregoing PCT and EP publications refer to the use of the compounds of formula I or II, and their pharmaceutically acceptable salts, in the treatment of illnesses induced or facilitated by CRF and in the treatment of anxiety, cardiovascular diseases, depression, fatigue syndrome, gastrointestinal diseases, headache, pain, cancer, immune dysfunction, hemorrhagic stress, drug addiction, drug and alcohol withdrawal symptoms, fertility problems, stress-induced psychotic episodes, neurodegenerative diseases such as Alzheimer's disease; irritable bowel syndrome 20 including Crohn's disease, spastic colon and irritable colon; eating disorders such as anorexia nervosa; and inflammatory disorders such as arthritis, asthma and allergies. Other CRF antagonists useful in the compositions and methods of this invention are referred to in commonly assigned PCT publication numbers WO 95/33727, WO 94/13644 and WO 94/13643. Each of the applications corresponding to PCT publication numbers WO 95/33727, WO 94/13644 and WO 25 94/13643 designate, inter alia, the United States and are incorporated herein by reference in their entirety.

[0005] RAS inhibitors useful in the practice of the instant invention include renin inhibitors, angiotensin-converting enzyme (ACE) inhibitors, and angiotensin-II (A-II) antagonists. Such RAS inhibitors are useful in the lowering of blood pressure in mammals and find utility in the treatment of congestive heart failure as well as other therapeutic effects. Specific RAS inhibitors useful in the practice of the instant invention are disclosed and referenced in detail hereinbelow.

30 [0006] Until the invention described herein, there was no report of the use of, or intent to use, a CRF antagonist together with a RAS inhibitor to achieve any therapeutic effect including the preferred treatment of congestive heart failure or hypertension.

[0007] Further, until this invention, there was no report of a pharmaceutical composition comprising a CRF antagonist together with a RAS inhibitor, nor of the use or intent to use such a composition in the treatment of any condition 35 including congestive heart failure or hypertension.

Summary of the Invention

40 [0008] This invention relates to compositions and methods useful in achieving therapeutic effects such as the treatment of congestive heart failure or hypertension, which compositions comprise preferably synergistic therapeutically effective amounts of a CRF antagonist, a RAS inhibitor and a pharmaceutically acceptable carrier, vehicle or diluent and which methods comprise administering to an animal, preferably a mammal including a human subject or a companion animal in need of such treatment, preferably synergistic therapeutically effective amounts of a CRF antagonist and a RAS inhibitor.

45 [0009] Preferably, the invention relates to the use of a CRF antagonist in combination with a RAS inhibitor selected from the group consisting of an angiotensin-converting enzyme (ACE) inhibitor, an angiotensin II antagonist and a renin inhibitor. More preferably, the renin angiotensin system inhibitor is a compound selected from the group consisting of benazeprilat, captopril, enalapril, lisinopril, losartan, valsartan, irbesartan, candesartan, telmisartan, tasosartan and eprosartan.

50 [0010] A preferred class of CRF antagonists comprises compounds of structural formula I or II, below,



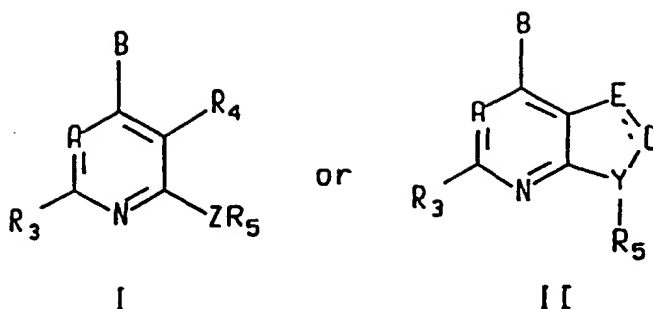
Detailed Description of the Invention

[0012] A first preferred aspect of the instant invention relates to compositions and methods for treating congestive heart failure, which compositions preferably comprise effective amounts, preferably synergistic, of a CRF antagonist, a RAS inhibitor and a pharmaceutically acceptable carrier, vehicle or diluent and which methods comprise administering to an animal, preferably a mammal including a human subject or a companion animal, such as a dog or cat, in need of such treatment, therapeutically effective amounts, preferably synergistic, of a CRF antagonist and a RAS inhibitor.

[0014] The term "synergistic" as utilized herein means that the effect achieved with the methods and compositions of the instant invention is greater than the sum of the effects that result from methods and compositions comprising the inhibitors and antagonists of this invention separately and in the amounts employed in the methods and compositions hereof.

[0016] The importance of CRF antagonists is set forth in the literature, e.g., as discussed in U.S. Patent 5,063,245, which is incorporated herein by reference, and a recent outline of the different activities possessed by CRF antagonists is found in Pharm. Rev., 43, 425-473 (1991). Such activity is readily determined by one skilled in the art according to standard assays including the methods described in Endocrinology, 116, 1653-1659 (1985) and Peptides, 10, 179-188 (1989) which determine the binding affinity of a test compound for a CRF receptor. The binding affinities for the active compounds, expressed as IC₅₀ values, generally range from about 0.2 nanomolar to about 10 micromolar. A variety of additional compounds having utility as CRF antagonists are, or will be, known to those skilled in the art. For example, CRF antagonists are mentioned in U.S. Patents 4,605,642 and 5,063,245 referring to certain peptides and pyrazolinones respectively, the teachings of which are incorporated herein by reference.

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15 including the pharmaceutically acceptable salts thereof, wherein the dashed line represents an optional double bond;

A is $-\text{CR}_7$ or N;

20 B is $-\text{NR}_1\text{R}_2$, $-\text{CR}_1\text{R}_2\text{R}_{11}$, $-\text{C}(=\text{CR}_1\text{R}_{12})\text{R}_2$, $-\text{NHC}\text{R}_{11}\text{R}_1\text{R}_2$, $-\text{OC}\text{R}_{11}\text{R}_1\text{R}_2$, $-\text{SC}\text{R}_{11}\text{R}_1\text{R}_2$, $-\text{CR}_{11}\text{R}_2\text{OR}_1$, $-\text{CR}_{11}\text{R}_2\text{SR}_1$, $-\text{C}(\text{S})\text{R}_2$, $-\text{NHN}\text{R}_1\text{R}_2$, $-\text{CR}_2\text{R}_{11}\text{NHR}_1$ or $-\text{C}(\text{O})\text{R}_2$;

D is: (i) N or $-\text{CR}_{10}$ when a double bond connects E and D and E is $-\text{CR}_4$; (ii) $-\text{CR}_{10}$ when a double bond connects E and D and E is N; (iii) $-\text{CR}_6\text{R}_9$, $-\text{CHR}_{10}$, $-\text{C}=\text{O}$, $-\text{C}=\text{S}$, $-\text{C}=\text{NH}$, or $-\text{C}=\text{NCH}_3$ when a single bond connects E and D; E is $-\text{CR}_4$ or N when a double bond connects E and D, and E is $-\text{CR}_4\text{R}_6$ or $-\text{NR}_6$ when a single bond connects E and D;

25 Y is N or $-\text{CH}$;

Z is NH, O, S, $-\text{N}(\text{C}_1\text{-C}_2 \text{ alkyl})$ or $-\text{CR}_{12}\text{R}_{13}$, wherein R_{12} and R_{13} are each, independently, hydrogen, trifluoromethyl or methyl, or one of R_{12} and R_{13} is cyano and the other is hydrogen or methyl;

30 R_1 is hydrogen or $\text{C}_1\text{-C}_6$ alkyl which is optionally substituted with one or two substituents independently selected from hydroxy, cyano, nitro, fluoro, chloro, bromo, iodo, CF_3 , $\text{C}_1\text{-C}_4$ alkoxy, $-\text{O}-\text{CO}-(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{O}-\text{CO}-\text{NH}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{O}-\text{CO}-\text{N}(\text{C}_1\text{-C}_4 \text{ alkyl})(\text{C}_1\text{-C}_2 \text{ alkyl})$, $-\text{NH}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{N}(\text{C}_1\text{-C}_2 \text{ alkyl})(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{S}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{N}(\text{C}_1\text{-C}_4 \text{ alkyl})\text{CO}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{NHCO}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{CO}_2(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{CONH}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{CON}(\text{C}_1\text{-C}_4 \text{ alkyl})(\text{C}_1\text{-C}_2 \text{ alkyl})$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfinyl}$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfonyl}$, and $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfanyl}$, and wherein said $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_4$ alkoxy and the $\text{C}_1\text{-C}_4$ alkyl moieties in the foregoing R_1 groups optionally contain one double or triple bond;

35 R_2 is $\text{C}_1\text{-C}_6$ alkyl, heteroaryl, aryl, $(\text{heteroaryl})\text{C}_1\text{-C}_4 \text{ alkyl}$ or $(\text{aryl})\text{C}_1\text{-C}_4 \text{ alkyl}$ wherein said aryl and the aryl moiety of said $(\text{aryl})\text{C}_1\text{-C}_4 \text{ alkyl}$ are selected from the group consisting of phenyl and naphthyl, and said heteroaryl and the heteroaryl moiety of said $(\text{heteroaryl})\text{C}_1\text{-C}_4 \text{ alkyl}$ is selected from the group consisting of thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, furanyl, benzofuranyl, benzothiazolyl, isothiazolyl, thiazolyl, benzisothiazolyl, benzisoxazolyl, benzimidazolyl, indolyl, and benzoxazolyl; or R^2 is $\text{C}_3\text{-C}_8$ cycloalkyl or $(\text{C}_3\text{-C}_8 \text{ cycloalkyl})\text{C}_1\text{-C}_6 \text{ alkyl}$, wherein one or two of the ring carbons of said cycloalkyl having at least 4 ring members and the cycloalkyl moiety of said $(\text{C}_3\text{-C}_8 \text{ cycloalkyl})\text{C}_1\text{-C}_6 \text{ alkyl}$ having at least 4 ring members is optionally replaced by an oxygen or sulfur atom or by $-\text{NR}_{14}$ wherein R_{14} is hydrogen or $\text{C}_1\text{-C}_4$ alkyl; and wherein each of the foregoing R_2 groups is optionally substituted by from one to three substituents independently selected from chloro, fluoro and $\text{C}_1\text{-C}_4$ alkyl, or by one substituent selected from bromo, iodo, cyano, nitro, $\text{C}_1\text{-C}_6$ alkoxy, $-\text{O}-\text{CO}-(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{O}-\text{CO}-\text{N}(\text{C}_1\text{-C}_4 \text{ alkyl})(\text{C}_1\text{-C}_2 \text{ alkyl})$, $-\text{CO}_2(\text{C}_1\text{-C}_4 \text{ alkyl})$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfanyl}$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfinyl}$, and $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfonyl}$, and wherein said $\text{C}_1\text{-C}_6$ alkyl and the $\text{C}_1\text{-C}_4$ alkyl and $\text{C}_1\text{-C}_6$ alkyl moieties of the foregoing R_2 groups optionally contain one carbon-carbon double or triple bond;

40 or R^1 and R^2 of said $-\text{NR}_1\text{R}_2$ and said $-\text{CR}_1\text{R}_2\text{R}_{11}$ are taken together to form a saturated 5 to 8 member ring, wherein said ring optionally contains one or two carbon-carbon double bonds, and wherein one or two of the ring carbons is optionally replaced by a heteroatom selected from O, S and N;

50 R_3 is hydrogen, $\text{C}_1\text{-C}_6$ alkyl, fluoro, chloro, bromo, iodo, hydroxy, amino, SH, $-\text{NH}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{N}(\text{C}_1\text{-C}_4 \text{ alkyl})(\text{C}_1\text{-C}_2 \text{ alkyl})$, $-\text{CH}_2\text{OH}$, $-\text{CH}_2\text{OCH}_3$, $-\text{O}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfanyl}$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfonyl}$, or $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfinyl}$, wherein said $\text{C}_1\text{-C}_6$ alkyl and the $\text{C}_1\text{-C}_4$ alkyl moieties of the foregoing R_3 groups optionally contain one double or triple bond and are optionally substituted by from one to three substituents independently selected from hydroxy, amino, $\text{C}_1\text{-C}_3$ alkoxy,

55 $-\text{NH}(\text{C}_1\text{-C}_2 \text{ alkyl})$, $-\text{N}(\text{C}_1\text{-C}_2 \text{ alkyl})_2$, $-\text{NHCOCH}_3$, fluoro, chloro and $\text{C}_1\text{-C}_3$ thioalkyl;

R_4 is hydrogen, $\text{C}_1\text{-C}_6$ alkyl, fluoro, chloro, bromo, iodo, $\text{C}_1\text{-C}_6$ alkoxy, formyl, trifluoromethoxy, $-\text{CH}_2\text{OCH}_3$, $-\text{CH}_2\text{OCH}_2\text{CH}_3$, $-\text{CH}_2\text{CH}_2\text{OCH}_3$, $-\text{CH}_2\text{CF}_3$, CF_3 , amino, nitro, $-\text{NH}(\text{C}_1\text{-C}_4 \text{ alkyl})$, $-\text{N}(\text{CH}_3)_2$, $-\text{NHCOCH}_3$, $-\text{NHCONHCH}_3$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfanyl}$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfinyl}$, $(\text{C}_1\text{-C}_4 \text{ alkyl})\text{sulfonyl}$, cyano, hydroxy, $-\text{CO}(\text{C}_1\text{-C}_4 \text{ alkyl})$,

- CHO, or -CO₂(C₁-C₄ alkyl), wherein said C₁-C₆ alkyl, C₁-C₆ alkoxy and the C₁-C₄ alkyl moieties of the foregoing R₄ groups optionally contain one double or triple bond and are optionally substituted with one substituent selected from hydroxy, amino, -NHCOCH₃, -NH(C₁-C₂ alkyl), -N(C₁-C₂ alkyl)₂, -CO₂(C₁-C₄ alkyl), -CO(C₁-C₄ alkyl), C₁-C₃ alkoxy, (C₁-C₃ alkyl)sulfanyl, fluoro, chloro, cyano and nitro;
- 5 R₅ is phenyl, naphthyl, thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, furanyl, benzofuranyl, benzothiazolyl, isothiazolyl, benzoisothiazolyl, thiazolyl, isoxazolyl, benzisoxazolyl, benzimidazolyl, triazolyl, pyrazolyl, pyrrolyl, indolyl, azaindolyl, benzoxazolyl, oxazolyl, pyrrolidinyl, thiazolidinyl, morpholinyl, pyridinyl, tetra-
- 10 zolyl, or 3- to 8-membered cycloalkyl or 9- to 12-membered bicycloalkyl, wherein said cycloalkyl and bicycloalkyl optionally contain one or two of O, S or -N-G wherein G is hydrogen, C₁-C₄ alkyl, C₁-C₄ alkanoyl, phenyl or benzyl, wherein each of the above R₅ groups is optionally substituted by from one to three substituents independently selected from fluoro, chloro, C₁-C₆ alkyl, C₁-C₆ alkoxy and trifluoromethyl, or one substituent selected from bromo, iodo, cyano, nitro, amino, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), -CO(C₁-C₄ alkyl), -
- 15 SO₂NH(C₁-C₄ alkyl), -SO₂N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -SO₂NH₂, -NHSO₂(C₁-C₄ alkyl), -S(C₁-C₄ alkyl), and -SO₂(C₁-C₄ alkyl), wherein said C₁-C₄ alkyl and C₁-C₆ alkyl moieties of the foregoing R₅ groups optionally contain one double or triple bond and are optionally substituted by one or two substituents independently selected from fluoro, chloro, hydroxy, amino, methylamino, dimethylamino and acetyl;
- R₆ is hydrogen or C₁-C₆ alkyl, wherein said C₁-C₆ alkyl is optionally substituted by a single hydroxy, methoxy, ethoxy or fluoro group;
- 20 R₇ is hydrogen, C₁-C₄ alkyl, fluoro, chloro, bromo, iodo, cyano, hydroxy, C₁-C₄ alkoxy, -CO(C₁-C₄ alkyl), -CO₂(C₁-C₄ alkyl), -OCF₃, CF₃, CH₂OH, -CH₂OCH₃ or -CH₂OCH₂CH₃;
- R₈ and R₉ are each, independently, hydrogen, hydroxy, methyl, ethyl, methoxy, or ethoxy; or R₈ and R₉ together form an oxo (=O) group;
- R₁₀ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, C₁-C₆ alkoxy, formyl, amino, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -SO_n(C₁-C₄ alkyl, wherein n is 0, 1 or 2, cyano, carboxy, or amido, wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl moieties of the foregoing R₁₀ groups are optionally substituted by one of hydroxy, trifluoromethyl,
- 25 amino, carboxy, amido, -NHCO(C₁-C₄ alkyl), -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), C₁-C₃ alkoxy, C₁-C₃ thioalkyl, fluoro, bromo, chloro, iodo, cyano or nitro; and
- R₁₁ is hydrogen, hydroxy, fluoro, or methoxy.
- 30 [0018] The term "alkyl", as used herein, unless otherwise indicated, includes saturated monovalent hydrocarbon radicals having straight or branched moieties or combinations thereof.
- [0019] The term "alkoxy", as used herein, unless otherwise indicated, includes O-alkyl groups wherein "alkyl" is defined above.
- [0020] The term "treatment", as used herein, unless otherwise indicated, includes the treatment, prevention, or inhibition of any disorder enumerated within the methods of the invention.
- 35 [0021] More specific compounds for use in the compositions and methods of the invention include compounds of formula I or II, or pharmaceutically acceptable salts thereof, wherein: B is -NR₁R₂, -NHCHR₁R₂, -CR₁R₂R₁₁, -SCHR₁R₂ or -OCHR₁R₂; R₁ is C₁-C₆ alkyl which is optionally substituted with a single hydroxy, fluoro or C₁-C₂ alkoxy group and optionally contains one carbon-carbon double or triple bond; R₂ is benzyl or C₁-C₆ alkyl which optionally contains one carbon-carbon double or triple bond, wherein said C₁-C₆ alkyl and the phenyl moiety of said benzyl are optionally substituted with fluoro, C₁-C₂ alkyl, or C₁-C₂ alkoxy; and R₁₁ is hydrogen or fluoro.
- 40 [0022] Other more specific compounds for use in the compositions and methods of the invention include compounds of formula I or II, or pharmaceutically acceptable salts thereof, wherein R₂ is (aryl)C₁-C₄ alkyl or (heteroaryl)C₁-C₄ alkyl in which said aryl or heteroaryl moiety is phenyl, thienyl, benzofuranyl, furanyl, benzothienyl, thiazolyl, pyridyl or benzothiazolyl.
- 45 [0023] Other more specific compounds for use in the compositions and methods of the invention include compounds of formula I or II, or pharmaceutically acceptable salts thereof, wherein B is -NR₁R₂ or -CHR₁R₂ in which R₁ and R₂ are taken together with N or CH to form a 5- or 6-membered ring optionally having sulfur, oxygen, or one more nitrogen atoms in said ring, such as a pyrrolidinyl, pyrrolyl, pyrazolyl, imidazolyl, oxazolyl, thiazolyl, isoxazolyl, thiadiazolyl, oxadiazolyl, pyridyl, pyrazinyl or pyrimidyl group.
- 50 [0024] Other more specific compounds for use in the compositions and methods of the invention include compounds of formula I or II, or pharmaceutically acceptable salts thereof, wherein B is -NHCHR₁R₂ or -OCHR₁R₂, wherein the CHR₁R₂ moiety is a 5- or 6-membered ring which optionally contains one oxygen or sulfur, such as a tetrahydrofuranyl, tetrahydrothiafuranyl or cyclopentanyl group.
- 55 [0025] Other more specific compounds for use in the compositions and methods of the invention include compounds of formula I or II, or pharmaceutically acceptable salts thereof, wherein B is tetrahydrofuranyl, tetrahydrothiafuranyl, tetrahydrothienyl, thiazolidinyl or cyclopentanyl.
- [0026] Other more specific compounds for use in the compositions and methods of the invention include com-

pounds of formula I or II, or pharmaceutically acceptable salts thereof, wherein R₃ is methyl, chloro, or methoxy; R₄ is methyl, -CH₂OH, cyano, trifluoromethoxy, methoxy, trifluoromethyl, chloro, -CO₂CH₃, -CH₂OCH₃, -CH₂Cl, -CH₂F, amino, nitro, hydrogen, methylsulfinyl, methylsulfonyl, methylsulfonyl, or ethyl; and R₅ is phenyl or pyridyl wherein said phenyl or pyridyl is substituted by one substituent independently selected from fluoro, chloro, bromo, iodo, C₁-C₄ alkoxy, trifluoromethyl, C₁-C₃ hydroxyalkyl, hydroxy, formyl, -CO₂(C₁-C₂ alkyl), (amino)C₁-C₂ alkyl, -CO(C₁-C₄ alkyl), and C₁-C₆ alkyl, wherein said C₁-C₆ alkyl and said C₁-C₄ alkyl are optionally substituted by a single hydroxy, or fluoro group and optionally contains one carbon-carbon double or triple bond.

[0027] For use in the compositions and methods of the invention, specific compounds of formula I or II include:

- 10 4-(1-ethyl-propoxy)-2,5-dimethyl-6-(2,4,6-trimethyl-benzyl)-pyrimidine;
2-(4-bromo-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
2-(4-ethyl-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
3-ethyl-4-(1-ethyl-propoxy)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
2-(2,6-dimethyl-4-propyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
- 15 4-(1-ethyl-propoxy)-2-(4-methoxy-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridine;
2-(4-ethoxy-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
2-(4-chloro-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
4-(1-methoxymethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
[3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-diethyl-amine;
- 20 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-propyl-amine;
[2,5-dimethyl-6-(2,4,6-trimethyl-phenoxy)-pyrimidin-4-yl](1-ethyl-propyl)-amine;
butyl-[3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-amine;
4-(1-ethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethyl-phenylsulfonyl)-pyridine;
butyl-[2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-ethyl-amine;
- 25 4-(1-ethyl-propylamino)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-nicotinic acid methyl ester,
[3,6-dimethyl-2-(2,4,6-trimethyl-phenylsulfonyl)-pyridin-4-yl]-ethyl-propyl-amine;
[4-(1-ethyl-propylamino)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-3-yl]-methanol;
[2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-ethyl-propyl-amine;
1-(ethyl-propyl)-[6-methyl-3-nitro-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-amine;
- 30 N4-(1-ethyl-propyl)-6-methyl-3-nitro-N2-(2,4,6-trimethyl-phenyl)-pyridine-2,4-diamine;
N4-(1-ethyl-propyl)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
N4-(1-ethyl-propyl)-6-methyl-N2-(2,4,6-trimethyl-phenyl)-pyridine-2,3,4-triamine;
[3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-(2,2,2-trifluoroethyl)-amine;
[3-chloromethyl-6-methyl-2-(2,4,6-trimethyl-phenoxy)pyridin-4-yl]-(1-ethyl-propyl)-amine;
- 35 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-(1-ethyl-propyl)-amine; (1-ethyl-propyl)-[2-methyl-5-nitro-6-(2,4,6-trimethyl-pyridin-3-yloxy)-pyrimidin-4-yl]-amine;
(1-ethyl-propyl)-[3-methoxymethyl-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-amine;
N-(1-ethyl-propyl)-2-methyl-5-nitro-N-(2,4,6-trimethyl-pyridin-3-yl)-pyrimidine-4,6-diamine;
[2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-diethyl-amine;
- 40 4-(1-ethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethylphenoxy)-pyridine;
butyl-[2,5-dimethyl-7-(2,4,6-trimethyl(phenyl)-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidin-4-yl]-ethyl-amine;
4-(butyl-ethylamino)-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-5,7-dihydro-pyrrolo[2,3-d]pyrimidin-6-one;
4-(1-ethylpropoxy)-2,5-dimethyl-6-(2,4,6-trimethylphenoxy)-pyrimidine;
N-butyl-N-ethyl-2,5-dimethyl-N-(2,4,6-trimethylphenyl)-pyrimidine-4,6-diamine;
- 45 (1-ethyl-propyl)-[5-methyl-3-(2,4,6-trimethyl-phenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-amine;
[2,5-dimethyl-3-(2,4,6-trimethyl-phenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-(1-ethyl-propyl)-amine;
N4-(1-ethyl-propyl)-6,N3,N3-trimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
N4-(1-ethyl-propyl)-6,N3,N3-trimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
6-(1-ethyl-propoxy)-2-methyl-N4-(2,4,6-trimethyl-phenyl)-pyrimidine-4,5-diamine;
- 50 [4-(1-ethyl-propoxy)-3,6-dimethyl-pyridin-2-yl]-(2,4,6-trimethylphenyl)-amine;
6-(ethyl-propyl-amino)-2,7-dimethyl-9-(2,4,6-trimethylphenyl)-7,9-dihydro-purin-8-one; and the pharmaceutically acceptable salts thereof.

[0028] For use in the compositions and methods of the invention, specific compounds of formula II wherein E and D are connected by a double bond, E is -CR₄, D is -CR₁₀ or N, Y is N, and A is -CR₇, include:

butyl-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-ethylamine;
3,6-dimethyl-4-(tetrahydrofuran-3-yloxy)-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridine;

[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-(1-methoxymethylpropyl)-amine;
 4-(1-methoxymethylpropoxy)-3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridine;
 (1-ethylpropyl)-[3,5,6-trimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-amine;
 4-(1-ethylpropoxy)-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-b]pyridine;
 4-(1-ethylpropoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-b]pyridine;
 4-(1-ethylpropoxy)-2,5-dimethyl-7-(2,6-dimethyl-4-bromophenyl)-7H-pyrrolo[2,3-b]pyridine; and the pharmaceutically acceptable salts thereof.

[0029] For use in the compositions and methods of the invention, specific compounds of formula II wherein E and D are connected by a double bond, E is -CR₄, and D, Y and A are N, include:

3-[(4-methyl-benzyl)-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amino]-propan-1-ol;
 diethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 2-[butyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amino]-ethanol;
 dibutyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-cyclopropylmethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 di-1-propyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 diallyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-chloro-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methoxy-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 propyl-ethyl-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 4-(1-ethyl-propyl)-6-methyl-3-methylsulfanyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidine;
 2-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino]-butan-1-ol;
 [3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-(1-methylpropyl)amine;
 4-(1-methoxymethylpropoxy)-3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidine; and the pharmaceutically acceptable salts thereof.

[0030] For use in the compositions and methods of the invention, specific compounds of formula II wherein E and D are connected by a double bond, E is -CR₄, D is -CR₁₀, and Y and A are N, include:

n-butyl-ethyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 di-n-propyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 ethyl-n-propyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 diethyl-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 n-butyl-ethyl-[2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 2-[N-n-butyl-N-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amino]-ethanol;
 4-(1-ethyl-propyl)-2,5-methyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 n-butyl-ethyl-[2,5-dimethyl-7-(2,4-dimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-(1-ethyl-propyl)amine;
 2-[7-(4-bromo-2,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-(S)-[7-(4-bromo-2,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 4-(1-ethyl-propoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 4-(1-methoxymethyl-propoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 4-(1-ethyl-butyl)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 [7-(4-bromo-2,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-(1-methoxymethyl-propyl)-amine;
 2-[7-(2-bromo-4,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(4-ethyl-2,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(2-ethyl-4,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(2-fluoromethyl-4,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol; and the pharmaceutically acceptable salts thereof.

[0031] The compounds of formula I or II, and their pharmaceutically acceptable salts, are readily prepared. The compounds of formula II wherein A, D and Y are N, a double bond connects E and D, and E is -CR₄, are prepared by one or more of the synthetic methods disclosed in PCT publication number WO 94/13677, referred to hereinabove and incorporated herein by reference. The compounds of formula II wherein A and Y are N, a double bond connects E and

D, E is -CR₄, and D is -CR₁₀, are prepared by one or more of the synthetic methods disclosed in PCT publication number WO 94/13676, referred to hereinabove and incorporated herein by reference. The compounds of formula II wherein A is -CR₇, a double bond connects E and D, E is -CR₄, D is N or -CR₁₀, and Y is N, are prepared by one or more of the synthetic methods disclosed in PCT publication number WO 95/34563, referred to hereinabove and incorporated herein by reference. The remaining compounds of formula II and the compounds of formula I are prepared by one or more of the synthetic methods disclosed in PCT publication number WO 95/33750, referred to hereinabove and incorporated herein by reference.

[0032] Pharmaceutically acceptable salts of the compounds of formula I and II include salts of acidic or basic groups. For example, pharmaceutically acceptable salts include sodium, calcium and potassium salts of acidic groups, such as when the R₁₀ substituent is carboxy. Such salts are generally prepared by combining a compound of formula I or II with one molar equivalent of a suitable Na⁺, K⁺, or Ca⁺² counterion in a suitable solvent. Pharmaceutically acceptable acid addition salts of basic groups, such as amino groups, are formed by reacting the base form of a compound of formula I or II with an appropriate conjugate acid. Pharmaceutically acceptable salts of basic groups include hydrochloride, hydrobromide, sulfate, hydrogen sulfate, phosphate, hydrogen phosphate, dihydrogen phosphate, acetate, succinate, citrate, tartrate, lactate, mandelate, benzenesulfonate (besylate), methanesulfonate (mesylate) and p-toluenesulfonate (tosylate) salts. When the salt is of a monobasic acid (e.g., the hydrochloride, the hydrobromide, the p-toluenesulfonate, the acetate), at least one molar equivalent and usually a molar excess of the acid is employed. However, when such salts as the sulfate, the hemisuccinate, the hydrogen phosphate or the phosphate are desired, the appropriate and exact chemical equivalents of the conjugate acid will generally be used. The free base and the acid are usually combined in a co-solvent from which the desired salt precipitates, or can be otherwise advantageously isolated by concentration or addition of a non-solvent.

[0033] In the practice of the instant invention, the CRF antagonist is preferably used in synergistic combination with a RAS inhibitor. The term renin-angiotensin system (RAS) inhibitor refers to a compound having the ability to down-regulate the conversion of the naturally-occurring plasma glycoprotein angiotensinogen into the N-terminal decapeptide angiotensin I or the subsequent conversion of angiotensin I to the octapeptide angiotensin II. While angiotensin I is known to possess minimal vasoactive properties, angiotensin II is a potent vasoconstrictor which mediates increased systemic vascular resistance with concomitant deleterious impact on various haemodynamic parameters in dysfunctional states. Thus, the renin-angiotensin system is known to be a causative factor in hypertension and congestive heart failure. For a discussion of the mechanisms of action and clinical utilities of various RAS inhibitors see, for example, Drugs, 28, 144-169 (1984) and the references cited therein.

[0034] In the practice of the instant invention, any RAS inhibitor may be employed. It is generally preferred, however, that the RAS inhibitor be a compound selected from the group consisting of a renin inhibitor, an angiotensin-converting enzyme (ACE) inhibitor, and an angiotensin II (A-II) antagonist. Especially preferred RAS inhibitors are selected from the group consisting of benazeprilat, captopril, enalapril, lisinopril, losartan, valsartan, irbesartan, candesartan, telmisartan, tasosartan and eprosartan.

[0035] The term renin inhibitor refers to a compound having the ability to inhibit the initial, rate-limiting step in the RAS cascade, i.e. the renin-mediated, proteolytic conversion of angiotensinogen into the N-terminal decapeptide angiotensin I, the penultimate precursor to angiotensin II. For a review of various renin inhibitors see, for example, Pharm. Res., 4, 364-374 (1987). A variety of renin inhibitors are, or will be, known to one skilled in the art including those described in U.S. Pat Nos. 4,814,342, 4,855,303, and 4,895,834, the teachings of which are incorporated herein by reference.

[0036] The term angiotensin-converting enzyme (ACE) inhibitor refers to a compound having the ability to inhibit the cleavage of the N-terminal decapeptide angiotensin I to the vasoactive octapeptide angiotensin II. For a review of various ACE inhibitors see, for example, Am. J. Cardiol. 66, 7D-13D (1990). A variety of ACE inhibitors are, or will be, known to one skilled in the art including those described in U.S. Pat Nos. 4,046,889 and 4,374,829, the teachings of which are incorporated herein by reference.

[0037] A preferred ACE inhibitor is benazeprilat: (3S)-3-[[[(1S)-1-carboxy-3-phenylpropyl]amino]-2,3,4,5-tetrahydro-2-oxo-1H-1-benzazepine-1-acetic acid and the variously substituted 1-carboxymethyl-3-(carboxymethylamino)-2,3,4,5-tetrahydro-1H-[1]benzaepin-2-ones related thereto which are disclosed in U.S. Patent No. 4,410,520, the disclosure of which is incorporated herein by reference.

[0038] Another preferred ACE inhibitor is captopril: 1-[(2S)-3-mercapto-2-methylpropionyl]-L-proline and the various derivatives and compounds related thereto which are disclosed in U.S. Patent No. 4,105,776, the disclosure of which is incorporated herein by reference.

[0039] Other preferred ACE inhibitors are enalapril: 1-[N-[(S)-1-carboxy-3-phenylpropyl]-L-alanyl]-L-proline-1'-ethyl ester and lisinopril: 1-[N²-[(S)-1-carboxy-3-phenylpropyl]-L-lysyl]-L-proline and the various carboxyalkyl dipeptide derivatives and compounds related thereto which are disclosed in U.S. Patent No. 4,374,829, the disclosure of which is incorporated herein by reference.

[0040] The term angiotensin II (A-II) antagonist refers to a compound having the ability to inhibit the vasoactive

effects of endogenous angiotensin II by competitive blockade at the angiotensin receptor sites located in vascular smooth muscle and within the adrenal gland. For a detailed review of angiotensin II receptors and the various antagonists thereof see, for example, Pharmacol. Rev. 45, 206-242 (1993). A variety of A-II antagonists are, or will be, known to one skilled in the art including those described in U.S. Pat. Nos. 4,355,040 and 4,880,804, EP 253310, EP 323841, and EP 324377, the teachings of which are incorporated herein by reference.

[0041] A preferred A-II antagonist is losartan: 2-butyl-4-chloro-1-[*p*-(*o*-1H-tetrazol-5-ylphenyl)benzyl]imidazole-5-methanol, monopotassium salt and the various substituted imidazole derivatives and pharmaceutically acceptable salts related thereto which are disclosed in U.S. Patent No. 5,138,069, the disclosure of which is incorporated herein by reference.

[0042] Another preferred A-II antagonist is valsartan: N-[(*p*-(*o*-1H-tetrazol-5-yl-phenyl)benzyl]-N-valeryl-L-valine and the various derivatives, pharmaceutically acceptable salts and compounds related thereto which are disclosed in U.S. Patent No. 5,399,578, the disclosure of which is incorporated herein by reference.

[0043] Another preferred A-II antagonist is irbesartan: 2-*n*-butyl-4-spirocyclopentane-1-(((2'-tetrazol-5-yl)biphenyl-4-yl)-2-imidazolyl-5-one and the various derivatives, pharmaceutically acceptable salts and compounds related thereto which are disclosed in U.S. Patent No's. 5,270,317 and 5,352,788, the disclosures of which are incorporated herein by reference.

[0044] Another preferred A-II antagonist is candesartan: 1-(cyclohexyloxycarbonyloxy)ethyl-2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]benzimidazole-7-carboxylate and the pharmaceutically acceptable salts thereof which are disclosed in U.S. Patent No. 5,196,444, the disclosure of which is incorporated herein by reference.

[0045] Another preferred A-II antagonist is telmisartan: 4'-[[4-methyl-6-(1-methyl-2-benzimidazolyl)-2-propyl-1-benzimidazolyl]methyl]-2-biphenylcarboxylic acid and the various derivatives and compounds related thereto which are disclosed in European Patent Application No. 0 502 314.

[0046] Yet another preferred A-II antagonist is tasosartan: 5,8-dihydro-2,4-dimethyl-8-[*p*-(*o*-1H-tetrazol-5-ylphenyl)benzyl]pyrido[2,3-*d*]pyrimidin-7(6*H*)-one and the pharmaceutically acceptable salts thereof which are disclosed in U.S. Patent No. 5,149,699, the disclosure of which is incorporated herein by reference.

[0047] Still another preferred A-II antagonist is eprosartan: (*E*)-2-butyl-1-(*p*-carboxybenzyl)- α -2-thenylimidazole-5-acrylic acid and the pharmaceutically acceptable salts thereof which are disclosed in U.S. Patent No. 5,185,351, the disclosure of which is incorporated herein by reference.

[0048] The administration of the CRF inhibitor and the RAS antagonist including the preferred synergistic embodiments thereof, according to this invention can be sequential in time or simultaneous with the simultaneous method being generally preferred. For sequential administration, the CRF inhibitor and the RAS antagonist can be administered in any order. It is generally preferred that such administration be oral. It is even more preferred that the administration be oral and simultaneous. However, if the subject being treated is unable to swallow, or oral absorption is otherwise impaired or undesirable, another route of administration such as suppositories, parenteral (such as subcutaneous, intravenous, intramuscular, intrasternal and infusion techniques), or topical administration will be appropriate. When the CRF inhibitor and the RAS antagonist are administered sequentially, the administration of each can be by the same method or by different methods.

[0049] The pharmaceutical compositions of this invention comprise amounts of a CRF antagonist and a RAS inhibitor. Preferred compositions comprise congestive heart failure-treating or hypertension-treating amounts, preferably synergistic, of a CRF antagonist and a RAS inhibitor. Especially preferred compositions of the instant invention comprise a CRF antagonist of structural formula I or II, or a pharmaceutically acceptable salt thereof, and a RAS inhibitor selected from the group consisting of a renin inhibitor, an angiotensin converting enzyme (ACE) inhibitor, and an angiotensin II (A-II) antagonist. Especially preferred RAS inhibitors are those selected from the group consisting of benazeprilat, captopril, enalapril, lisinopril, losartan, valsartan, irbesartan, candesartan, telmisartan, tasosartan and eprosartan. It is further preferred that the composition comprising the CRF antagonist and the RAS inhibitor be administered in the presence of a pharmaceutically-acceptable carrier, vehicle or diluent. Suitable pharmaceutical carriers, vehicles and diluents useful in the administration of the above combinations include inert solid diluents or fillers, sterile aqueous solution and various organic solvents. The pharmaceutical compositions formed by combining the active compounds and the pharmaceutically acceptable carriers, vehicles or diluents are then readily administered in a variety of dosage forms such as tablets, powders, lozenges, syrups, injectable solutions and the like. These pharmaceutical compositions can, if desired, contain additional ingredients such as flavorings, binders, excipients and the like. Thus, for purposes of oral administration, tablets containing various excipients such as sodium citrate, calcium carbonate and calcium phosphate may be employed along with various disintegrants such as starch, alginic acid and certain complex silicates, together with binding agents such as polyvinylpyrrolidone, sucrose, gelatin and acacia. Additionally, lubricating agents such as magnesium stearate, sodium lauryl sulfate and talc are often useful for tableting purposes. Solid compositions of a similar type can also be employed as fillers in soft and hard filled gelatin capsules. Preferred materials for this include lactose or milk sugar and high molecular weight polyethylene glycols. When aqueous suspensions or elixirs are desired for oral administration, the essential active ingredient(s) therein may be combined with various sweetening

or flavoring agents, coloring matter or dyes and, if desired, emulsifying or suspending agents, together with diluents such as water, ethanol, propylene glycol, glycerin and combinations thereof, all of which are pharmaceutically acceptable.

[0050] For parenteral administration, solutions of the active compound(s) in sesame or peanut oil, aqueous propylene glycol, or in sterile aqueous solution can be employed. Such aqueous solutions should be suitably buffered if necessary and the liquid diluent first rendered isotonic with sufficient saline or glucose. These particular aqueous solutions are especially suitable for intravenous, intramuscular, subcutaneous and intraperitoneal administration. The sterile aqueous media employed are all readily available by standard techniques known to those skilled in the art.

[0051] For purposes of transdermal (e.g., topical) administration, dilute, sterile, aqueous or partially aqueous solutions (usually in about 0.1% to 5% concentration), otherwise similar to the above parenteral solutions, are employed.

[0052] Methods of preparing various pharmaceutical compositions with a certain amount of active ingredient are known, or will be apparent in light of this disclosure, to one skilled in the art. For example, see Remington's Pharmaceutical Sciences, Mack Publishing Company, Easton, Pa., 15th Edition (1975).

[0053] The dosage(s) of the CRF antagonist and the RAS inhibitor necessary to achieve the desired therapeutic effect according to this invention are within the skill of those who practice in the art of having the benefit of the disclosure herein.

[0054] Dosage ranges for certain CRF antagonists have been reported with representative dosages ranging from about 0.1 to 500 mg/kg of the body weight of the patient to be treated. However, some variability in the general dosage range may be required depending upon the age and weight of the patient, the intended route of administration, and the progress and degree of severity of the illness to be treated.

[0055] Dosage ranges for certain renin inhibitors have been reported with representative dosages being 0.250 mg/kg to 1.4 mg/kg i.v. and 40 mg/day to 1200 mg/day orally. Dosage ranges for certain ACE inhibitors have been reported with representative dosages of 50 mg/day to 450 mg/day orally and 20 mg/day parenterally. Dosage ranges for certain angiotensin II antagonists have been reported with representative dosages being about 0.5 to 500 mg/kg p.o., preferably 2 to 80 mg/kg p.o., and 3 mg/kg i.v. The dosages to be employed according to this invention may also be varied depending upon the requirements of the subject being treated, the degree of severity of the condition being treated and the compound being administered.

[0056] The methods and compositions of the instant invention also have utility in the treatment of congestive heart failure or hypertension in companion animals such as dogs and cats. The administration of the compositions of this invention may be effected orally or parenterally. An amount of a preferred synergistic composition of the invention is administered such that an effective dose is received, usually a daily dose. Conveniently, the medicaments can be carried in the drinking water such that a therapeutic dosage of the agents is ingested with the daily water supply. The agents can be directly metered into drinking water, preferably in the form of a liquid, water-soluble concentrate, such as an aqueous solution of a water-soluble salt.

[0057] Conveniently, the active ingredients can also be added directly to the companion animal's feed, as such, or in the form of an animal feed supplement, also referred to as a premix or concentrate. A premix or concentrate of therapeutic agent in a carrier is more commonly employed for the inclusion of the agent in the feed. Suitable carriers are liquid or solid, as desired, such as water, various meals such as alfalfa meal, soybean meal, cottonseed oil meal, linseed oil meal, corncob meal and corn meal, molasses, urea, bone meal, and various mineral mixes. A particularly effective carrier is the respective animal feed itself, i.e., a small portion of such feed. The carrier facilitates uniform distribution of the active materials in the finished feed with which the premix is blended. It is important that the compounds be thoroughly blended into the premix and, subsequently, the feed. In this respect, the agents may be dispersed or dissolved in a suitable oily vehicle such as soybean oil, corn oil, cottonseed oil, and the like, or in a volatile organic solvent and then blended with the carrier. It will be appreciated that the proportions of active materials in the concentrate are capable of wide variation since the amount of agent in the finished feed may be adjusted by blending the appropriate proportion of premix with the feed to obtain a desired level of the therapeutic agents.

[0058] High potency concentrates may be blended by the feed manufacturer with a proteinaceous carrier such as soybean oil meal and other meals, as described above, to produce concentrated supplements which are suitable for direct feeding to animals. In such instances, the animals are permitted to consume the usual diet. Alternatively, such concentrated supplements may be added directly to the feed to produce a nutritionally balanced, finished feed containing a therapeutically effective level of compound(s) according to the invention. The mixtures are thoroughly blended by standard procedures, such as in a twin shell blender, to insure homogeneity.

[0059] If the supplement is used as a top dressing for the feed, it likewise helps to insure uniformity of distribution of the active ingredient across the top of the dressed feed.

[0060] For veterinary uses, both paste and pellet formulations may also be conveniently employed. Paste formulations can be prepared readily by dispersing the active compounds in a pharmaceutically acceptable oil such as peanut oil, sesame oil, corn oil, and the like. Similarly, pellets containing an effective amount of the compounds of the instant invention can be prepared by admixing the compounds of the invention with a suitable diluent such as carbowax, car-

nuba wax, and the like, and a lubricant, such as magnesium or calcium stearate, can be employed to improve the pelleting process.

[0061] Since the instant invention relates to the treatment of congestive heart failure or hypertension with a combination, preferably synergistic, of active ingredients which may be administered separately, the invention also relates to combining separate pharmaceutical compositions in kit form. A kit, according to this invention, comprises two separate pharmaceutical compositions: a first unit dosage form, comprising a preferably synergistic amount of a CRF antagonist and a pharmaceutically acceptable carrier, vehicle or diluent and a second unit dosage form comprising a preferably synergistic amount of a RAS inhibitor and a pharmaceutically acceptable carrier, vehicle or diluent. The kit further comprises a container. The container is used to contain the separate compositions and may comprise, for example, a divided bottle or a divided foil packet however, the separate compositions may also be contained within a single, undivided container. Normally, the kit will also include directions for the administration of the separate components. The kit form is particularly advantageous when the separate components are preferably administered in different dosage forms (e.g., oral and parenteral), are administered at different dosage levels, or when titration of the individual components of the combination is desired by the prescribing physician.

[0062] An example of such a kit is a so-called blister pack. Blister packs are well known in the packaging industry and are being used widely for the packaging of pharmaceutical unit dosage forms (tablets, capsules and the like). Blister packs generally consist of a sheet of relatively rigid material covered with a foil of a preferably transparent plastic material. During the packaging process recesses are formed in the plastic foil. The recesses generally conform to the size and shape of the tablets or capsules to be contained therein. Next, the tablets or capsules are placed in the recesses and the sheet of relatively rigid material is sealed against the plastic foil at the face of the foil which is opposite from the direction in which the recesses were formed. As a result, the tablets or capsules are sealed in the recesses between the plastic foil and the sheet. Preferably, the strength of the sheet is such that the tablets or capsules may be removed from the blister pack by the application of manual pressure on the recesses whereby an opening is formed in the sheet at the place of the recess. The tablet or capsule can then be removed through the formed opening.

[0063] It is further desirable to provide a memory aid on the pack, e.g., in the form of numbers or similar indicia next to the tablets or capsules whereby the indicia correspond with the days of the regimen which the dosage form so specified is to be ingested. An additional example of such a memory aid is a calendar printed on the pack, e.g., as follows First Week, Monday, Tuesday, . . . etc. . . . Second Week, Monday, Tuesday, . . . " etc. Other variations will be readily apparent. A "daily dose" can be a single tablet or capsule or multiple tablets or capsules to be ingested on a given day. Also, a daily dose of a CRF antagonist can consist of one tablet or capsule while a daily dose of a RAS inhibitor can consist of multiple tablets or capsules, or vice versa. The memory aid should reflect this.

[0064] In another specific embodiment of the invention, a pack designed to dispense the daily doses one at a time in the order of their intended use is provided. Preferably, the pack is equipped with a memory aid, so as to further facilitate compliance with the dosage regimen. An example of such a memory aid is a mechanical counter which indicates the number of daily doses to be dispensed. Another example of such a memory aid is a battery-powered micro-chip memory coupled with a liquid crystal readout, or audible reminder signal which, for example, reads out the date that the last daily dose has been taken and/or reminds the patient when the next dose is to be taken.

[0065] Evaluation of the synergistic CRF antagonist - RAS inhibitor combination therapy for treating congestive heart failure and hypertension is performed utilizing the methodology of Sladek, et al., Cardiovascular Research, 31, 568-576 (1996).

[0066] Adult male Sprague-Dawley rats are anesthetized by intraperitoneal injection of ketamine and sodium pentobarbital. The heart is surgically exposed and the left coronary artery, in experimentally infarcted animals, is occluded by ligation. In sham-operated animals the ligature remains loose. The animals are allowed to recover for a period of three weeks.

[0067] Prior to surgery, the treated animals receive a pre-determined amount of the test compound and this dosing is continued throughout the recovery period. Three weeks after surgery, the animals from all experimental groups are anesthetized as described previously. Blood pressure in the left (LV) and right ventricles (RV) is measured using a catheter-tip pressure transducer. The catheter is inserted into the left ventricular chamber via the right carotid artery under continuous pressure monitoring. Determinations are recorded after a 15 minute stabilization period. Similarly, a catheter is introduced into the RV via the right jugular vein.

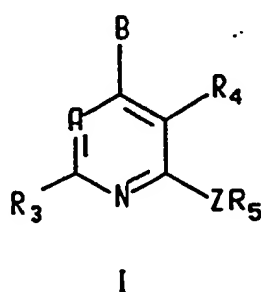
[0068] The analog pressure signal is digitized with a sampling frequency of 1 kHz and stored on a computer for later processing. The following parameters are derived: systolic pressure (LVSP, RVSP), left ventricular end-diastolic pressure (LVEDP), developed pressure (LVDevP) and the maximal rates of pressure development $(+dP/dt)_{\max}$ and fall $(-dP/dt)_{\max}$. In addition, the time constant of relaxation (τ) is calculated on the basis of an experimental model of isovolumetric pressure decay, as the time required for the pressure at $(+dP/dt)_{\max}$ to be reduced by $1/e$ (Mirsky, et al., Prog. Cardiovasc. Dis., 32, 291-318, (1990)). Heart rate is calculated from the left ventricular pressure signal.

[0069] The use of this method for assessment of synergy in the treatment of congestive heart failure or hypertension is deemed appropriate where a quantitative dose-response curve for each drug exists. In this instance, a synergis-

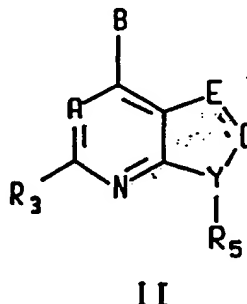
tic response is greater than an additive quantitative response obtained with a combination of two agents compared to the same biological response at a particular dose based on the single agent dose-response curve.

Claims

1. A method of achieving a therapeutic effect in an animal which method comprises administering to an animal therapeutically effective amounts of a corticotropin releasing factor antagonist and a renin-angiotensin system inhibitor.
2. A method of treating congestive heart failure or hypertension which method comprises administering to an animal in need of such treatment therapeutically effective amounts of a corticotropin releasing factor antagonist and a renin-angiotensin system inhibitor.
3. A method of claim 2 wherein said corticotropin releasing factor antagonist and said renin-angiotensin system inhibitor are present in synergistic therapeutically effective amounts.
4. A method of claim 2 wherein said renin-angiotensin system inhibitor is selected from the group consisting of a renin inhibitor, an angiotensin-converting enzyme inhibitor, and an angiotensin II antagonist
5. A method of claim 4 wherein said renin-angiotensin system inhibitor is selected from the group consisting of benazeprilat, captopril, enalapril, lisinopril, losartan, valsartan, irbesartan, candesartan, telmisartan, tasosartan and eprosartan.
6. A method of claim 2 wherein said corticotropin releasing factor antagonist is a compound of structural formula



or



or a pharmaceutically acceptable salt thereof, wherein the dashed line represents an optional double bond;

A is -CR₇ or N;

B is -NR₁R₂, -CR₁R₂R₁₁, -C(=CR₁R₁₂)R₂, -NHCR₁₁R₁R₂, -OCR₁₁R₁R₂, -SCR₁₁R₁R₂, -CR₁₁R₂OR₁, -CR₁₁R₂SR₁, -C(S)R₂, -NHNH₁R₂, -CR₂R₁₁NHR₁ or -C(O)R₂;

D is: (i) N or -CR₁₀ when a double bond connects E and D and E is -CR₄; (ii) -CR₁₀ when a double bond connects E and D and E is N; or (iii) -CR₈R₉, -CHR₁₀, -C=O, -C=S, -C=NH, or -C=NCH₃ when a single bond connects E and D;

E is -CR₄ or N when a double bond connects E and D, and E is -CR₄R₆ or -NR₆ when a single bond connects E and D;

Y is N or -CH;

Z is NH, O, S, -N(C₁-C₂ alkyl) or -CR₁₂R₁₃, wherein R₁₂ and R₁₃ are each, independently, hydrogen, trifluoromethyl or methyl, or one of R₁₂ and R₁₃ is cyano and the other is hydrogen or methyl;

R₁ is hydrogen or C₁-C₆ alkyl which is optionally substituted with one or two substituents independently selected from hydroxy, cyano, nitro, fluoro, chloro, bromo, iodo, CF₃, C₁-C₄ alkoxy, -O-CO-(C₁-C₄ alkyl), -O-CO-NH(C₁-C₄ alkyl), -O-CO-N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -NH(C₁-C₄ alkyl), -N(C₁-C₂ alkyl)(C₁-C₄ alkyl), -S(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)CO(C₁-C₄ alkyl), -NHCO(C₁-C₄ alkyl), -CO₂(C₁-C₄ alkyl), -CONH(C₁-C₄ alkyl), -CON(C₁-C₄ alkyl)(C₁-C₂ alkyl), (C₁-C₄ alkyl)sulfinyl, (C₁-C₄ alkyl)sulfonyl, and (C₁-C₄ alkyl)sulfanyl, and wherein said C₁-C₆ alkyl, C₁-C₄ alkoxy and the C₁-C₄ alkyl moieties in the foregoing R₁ groups optionally con-

tain one double or triple bond;

R₂ is C₁-C₆ alkyl, heteroaryl, aryl (heteroaryl)C₁-C₄ alkyl or (aryl)C₁-C₄ alkyl wherein said aryl and the aryl moiety of said (aryl)C₁-C₄ alkyl are selected from the group consisting of phenyl and naphthyl, and said heteroaryl and the heteroaryl moiety of said (heteroaryl)C₁-C₄ alkyl is selected from the group consisting of thienyl, benzothienyl, pyridyl, thiazolyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, furanyl, benzofuranyl, benzothiazolyl, isothiazolyl, benzisothiazolyl, benzisoxazolyl, benzimidazolyl, indolyl, and benzoxazolyl; or R² is C₃-C₈ cycloalkyl or (C₃-C₈ cycloalkyl)C₁-C₆ alkyl, wherein one or two of the ring carbons of said cycloalkyl having at least 4 ring members and the cycloalkyl moiety of said (C₃-C₈ cycloalkyl)C₁-C₆ alkyl having at least 4 ring members is optionally replaced with an oxygen or sulfur atom or with -NR₁₄ wherein R₁₄ is hydrogen or C₁-C₄ alkyl; and wherein each of the foregoing R₂ groups is optionally substituted with from one to three substituents independently selected from chloro, fluoro and C₁-C₄ alkyl, or with one substituent selected from bromo, iodo, cyano, nitro, C₁-C₆ alkoxy, -O-CO-(C₁-C₄ alkyl), -O-CO-N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), (C₁-C₄ alkyl)sulfanyl, (C₁-C₄ alkyl)sulfinyl, and (C₁-C₄ alkyl)sulfonyl, and wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl and C₁-C₆ alkyl moieties of the foregoing R₂ groups optionally contain one carbon-carbon double or triple bond;

or R¹ and R² of said -NR₁R₂ and said -CR₁R₂R₁₁ are taken together to form a saturated 5 to 8 member ring, wherein said ring optionally contains one or two carbon-carbon double bonds, and wherein one or two of the ring carbons is optionally replaced with a heteroatom selected from O, S and N;

R₃ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, hydroxy, amino, SH, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CH₂OH, -CH₂OCH₃, -O(C₁-C₄ alkyl), (C₁-C₄ alkyl)sulfanyl, (C₁-C₄ alkyl)sulfonyl, or (C₁-C₄ alkyl)sulfinyl, wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl moieties of the foregoing R₃ groups optionally contain one double or triple bond and are optionally substituted with from one to three substituents independently selected from hydroxy, amino, C₁-C₃ alkoxy,

-NH(C₁-C₂ alkyl), -N(C₁-C₂)₂, -NHCOCH₃, fluoro, chloro and C₁-C₃ thioalkyl;

R₄ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, C₁-C₆ alkoxy, formyl, trifluoromethoxy, -CH₂OCH₃, -CH₂OCH₂CH₃, -CH₂CH₂OCH₃, -CH₂CF₃, CF₃, amino, nitro, -NH(C₁-C₄ alkyl), -N(CH₃)₂, -NHCOCH₃, -NHCONHCH₃, (C₁-C₄ alkyl)sulfanyl, (C₁-C₄ alkyl)sulfinyl, (C₁-C₄ alkyl)sulfonyl, cyano, hydroxy, -CO(C₁-C₄ alkyl), -CHO, or -CO₂(C₁-C₄ alkyl), wherein said C₁-C₆ alkyl, C₁-C₆ alkoxy and the C₁-C₄ alkyl moieties of the foregoing R₄ groups optionally contain one double or triple bond and are optionally substituted with one substituent selected from hydroxy, amino, -NHCOCH₃, -NH(C₁-C₂ alkyl), -N(C₁-C₂ alkyl)₂, -CO₂(C₁-C₄ alkyl), -CO(C₁-C₄ alkyl), C₁-C₃ alkoxy, (C₁-C₃ alkyl)sulfanyl, fluoro, chloro, cyano and nitro;

R₅ is phenyl, naphthyl, thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, furanyl, benzofuranyl, benzothiazolyl, isothiazolyl, benzisothiazolyl, thiazolyl, isoxazolyl, benzisoxazolyl, benzimidazolyl, triazolyl, pyrazolyl, pyrrolyl, indolyl, azaindolyl, benzoxazolyl, oxazolyl, pyrrolidinyl, thiazolidinyl, morpholinyl, pyridinyl, tetrazolyl, or 3- to 8-membered cycloalkyl or 9- to 12-membered bicycloalkyl, wherein said cycloalkyl and bicycloalkyl optionally contain one or two of O, S or -N-G wherein G is hydrogen, C₁-C₄ alkyl, C₁-C₄ alkanoyl, phenyl or benzyl, wherein each of the above R₅ groups is optionally substituted with from one to three substituents independently selected from fluoro, chloro, C₁-C₆ alkyl, C₁-C₆ alkoxy and trifluoromethyl, or one substituent selected from bromo, iodo, cyano, nitro, amino, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), -CO(C₁-C₄ alkyl), -SO₂NH(C₁-C₄ alkyl), -SO₂N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -SO₂NH₂, -NHSO₂(C₁-C₄ alkyl), -S(C₁-C₄ alkyl), and -SO₂(C₁-C₄ alkyl), wherein said C₁-C₄ alkyl and C₁-C₆ alkyl moieties of the foregoing R₅ groups optionally contain one double or triple bond and are optionally substituted with one or two substituents independently selected from fluoro, chloro, hydroxy, amino, methylamino, dimethylamino and acetyl;

R₆ is hydrogen or C₁-C₆ alkyl, wherein said C₁-C₆ alkyl is optionally substituted with a single hydroxy, methoxy, ethoxy or fluoro group;

R₇ is hydrogen, C₁-C₄ alkyl, fluoro, chloro, bromo, iodo, cyano, hydroxy, C₁-C₄ alkoxy, -CO(C₁-C₄ alkyl), -CO₂(C₁-C₄ alkyl), -OCF₃, CF₃, -CH₂OH, -CH₂OCH₃ or -CH₂OCH₂CH₃;

R₈ and R₉ are each, independently, hydrogen, hydroxy, methyl, ethyl, methoxy, or ethoxy;

or R₈ and R₉ together form an oxo (=O) group;

R₁₀ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, C₁-C₆ alkoxy, formyl, amino, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -SO_n(C₁-C₄ alkyl), wherein n is 0, 1 or 2, cyano, carboxy, or amido, wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl moieties of the foregoing R₁₀ groups are optionally substituted with one of hydroxy, trifluoromethyl, amino, carboxy, amido, -NHCO(C₁-C₄ alkyl), -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), C₁-C₃ alkoxy, C₁-C₃ thioalkyl, fluoro, bromo, chloro, iodo, cyano or nitro; and,

R₁₁ is hydrogen, hydroxy, fluoro, or methoxy.

7. A method of claim 6 wherein B is -NR₁R₂, -NHCHR₁R₂, -CR₁R₂R₁₁, -SCHR₁R₂ or -OCHR₁R₂; R₁ is C₁-C₆ alkyl

which is optionally substituted with a single hydroxy, fluoro or C₁-C₂ alkoxy group and optionally contains one carbon-carbon double or triple bond; R₂ is benzyl or C₁-C₆ alkyl which optionally contains one carbon-carbon double or triple bond, wherein said C₁-C₆ alkyl and the phenyl moiety of said benzyl are optionally substituted with fluoro, C₁-C₂ alkyl, or C₁-C₂ alkoxy; and R₁₁ is hydrogen or fluoro.

8. A method of claim 6 wherein R₂ is (aryl)C₁-C₄ alkyl or (heteroaryl)C₁-C₄ alkyl in which said aryl or heteroaryl moiety is phenyl, thienyl, benzofuranyl, furanyl, benzothienyl, thiazolyl, pyridyl or benzothiazolyl.
9. A method of claim 6 wherein B is NR₁R₂ or CHR₁R₂ in which R₁ and R₂ are taken together with N or CH to form a 5- or 6-membered ring optionally having sulfur, oxygen, or, where B is NR₁R₂, one more nitrogen in said ring.
10. A method of claim 6 wherein B is -NHCHR₁R₂ or -OCHR₁R₂, wherein the CHR₁R₂ moiety is a 5- or 6-membered ring which optionally contains one oxygen or sulfur.
11. A method of claim 10 wherein B is tetrahydrofuranyl, tetrahydrothiafuranyl, tetrahydrothienyl, thiazolidinyl or cyclopentanyl.
12. A method of claim 6 wherein R₃ is methyl, chloro, or methoxy; R₄ is methyl, -CH₂OH, cyano, trifluoromethoxy, methoxy, trifluoromethyl, chloro, -CO₂CH₃, -CH₂OCH₃, -CH₂Cl, -CH₂F, amino or nitro; R₆ is hydrogen, methylsulfinyl, methylsulfonyl, methylsulfanyl, or ethyl; and R₅ is phenyl or pyridyl wherein said phenyl or pyridyl is substituted with one substituent independently selected from fluoro, chloro, bromo, iodo, C₁-C₄ alkoxy, trifluoromethyl, C₁-C₃ hydroxyalkyl, -CO₂(C₁-C₂ alkyl), (amino)C₁-C₂ alkyl, -CO(C₁-C₄ alkyl), and C₁-C₆ alkyl, wherein said C₁-C₆ alkyl and said C₁-C₄ alkyl are optionally substituted with a single hydroxy, or fluoro group and optionally contains one carbon-carbon double or triple bond.
13. A method of claim 6 wherein the compound of formula I or II, is selected from the group consisting of:

4-(1-ethyl-propoxy)-2,5-dimethyl-6-(2,4,6-trimethyl-benzyl)-pyrimidine;
 2-(4-bromo-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 2-(4-ethyl-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 3-ethyl-4-(1-ethyl-propoxy)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
 2-(2,6-dimethyl-4-propyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 4-(1-ethyl-propoxy)-2-(4-methoxy-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridine;
 2-(4-ethoxy-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 2-(4-chloro-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 4-(1-methoxymethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-diethyl-amine;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-propyl-amine;
 [2,5-dimethyl-6-(2,4,6-trimethyl-phenoxy)-pyrrolidin-4-yl]-(1-ethyl-propyl)-amine;
 butyl-[3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-amine;
 4-(1-ethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethyl-phenylsulfanyl)-pyridine;
 butyl-[2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-ethyl-amine;
 4-(1-ethyl-propylamino)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-nicotinic acid methyl ester;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenylsulfanyl)-pyridin-4-yl]-ethyl-propyl-amine;
 [4-(1-ethyl-propylamino)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-3-yl]-methanol;
 [2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-ethyl-propyl-amine;
 1-(ethyl-propyl)-[6-methyl-3-nitro-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-amine;
 N4-(1-ethyl-propyl)-6-methyl-3-nitro-N2-(2,4,6-trimethyl-phenyl)-pyridine-2,4-diamine;
 N4-(1-ethyl-propyl)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
 N4-(1-ethyl-propyl)-6-methyl-N2-(2,4,6-trimethyl-phenyl)-pyridine-2,3,4-triamine;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-(2,2,2-trifluoro-ethyl)-amine;
 [3-chloromethyl-6-methyl-2-(2,4,6-trimethyl-phenoxy)pyridin-4-yl]-(1-ethyl-propyl)-amine;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-(1-ethyl-propyl)-amine;
 (1-ethyl-propyl)-[2-methyl-5-nitro-6-(2,4,6-trimethyl-pyridin-3-yloxy)-pyrimidin-4-yl]-amine;
 (1-ethyl-propyl)-[3-methoxymethyl-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-amine;
 N-(1-ethyl-propyl)-2-methyl-5-nitro-N-(2,4,6-trimethyl-pyridin-3-yl)-pyrimidine-4,6-diamine;
 [2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-diethyl-amine;
 4-(1-ethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethylphenoxy)-pyridine;

butyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidin-4-yl]-ethyl-amine;
 4-(butyl-ethylamino)-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-5,7-dihydro-pyrrolo[2,3-d]pyrimidin-6-one;
 4-(1-ethylpropoxy)-2,5-dimethyl-6-(2,4,6-trimethylphenoxy)-pyrimidine;
 N-butyl-N-ethyl-2,5-dimethyl-N-(2,4,6-trimethylphenyl)-pyrimidine-4,6-diamine;
 5 (1-ethyl-propyl)-[5-methyl-3-(2,4,6-trimethyl-phenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-amine;
 [2,5-dimethyl-3-(2,4,6-trimethyl-phenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-(1-ethyl-propyl)-amine;
 N4-(1-ethyl-propyl)-6,N3-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
 N4-(1-ethyl-propyl)-6,N3,N3-trimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
 6-(1-ethyl-propoxy)-2-methyl-N4-(2,4,6-trimethyl-phenyl)-pyrimidine-4,5-diamine;
 10 [4-(1-ethyl-propoxy)-3,6-dimethyl-pyridin-2-yl]-(2,4,6-trimethylphenyl)-amine;
 6-(ethyl-propyl-amino)-2,7-dimethyl-9-(2,4,6-trimethylphenyl)-7,9-dihydro-purin-8-one; and the pharmaceuti-
 cally acceptable salts thereof.

14. A method of claim 6 wherein said compound is a compound of formula II in which E and D are connected by a dou-
 15 ble bond, E is -CR₄, D is -CR₁₀ or N, Y is N, and A is -CR₇, or a pharmaceutically acceptable salt thereof.

15. A method of claim 14 wherein said compound is selected from the group consisting of:

butyl-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-ethylamine;
 20 3,6-dimethyl-4-(tetrahydrofuran-3-yloxy)-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridine
 [3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-(1-methoxymethylpropyl)-amine;
 4-(1-methoxymethylpropoxy)-3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridine;
 (1-ethylpropyl)-[3,5,6-trimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-amine;
 4-(1-ethylpropoxy)-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-b]pyridine;
 25 4-(1-ethylpropoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-b]pyridine;
 4-(1-ethylpropoxy)-2,5-dimethyl-7-(2,6-dimethyl-4-bromophenyl)-7H-pyrrolo[2,3-b]pyridine; and the pharma-
 ceutically acceptable salts thereof.

16. A method of claim 6 wherein said compound is a compound of formula II in which E and D are connected by a dou-
 30 ble bond, E is -CR₄, and D, Y and A are N, or a pharmaceutically acceptable salt thereof.

17. A method of claim 16 wherein said compound is selected from the group consisting of:

3-(4-methyl-benzyl)-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amino}-propan-
 35 1-ol;
 diethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 2-butyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amino}-ethanol;
 dibutyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 40 butyl-ethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-cyclopropylmethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-
 yl]-amine;
 di-1-propyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 diallyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 45 butyl-ethyl-[6-chloro-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methoxy-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 propyl-ethyl-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 4-(1-ethyl-propyl)-6-methyl-3-methylsulfanyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidine;
 2-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamine]-butan-1-ol;
 50 [3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-(1-methylpropyl)amine;
 4-(1-methoxymethylpropoxy)-3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidine; and the
 pharmaceutically acceptable salts thereof.

18. A method of claim 6 wherein said compound is a compound of formula II in which E and D are connected by a dou-
 55 ble bond, E is -CR₄, D is -CR₁₀, and Y and A are N, or a pharmaceutically acceptable salt thereof.

19. A method of claim 18 wherein said compound is selected from the group consisting of:

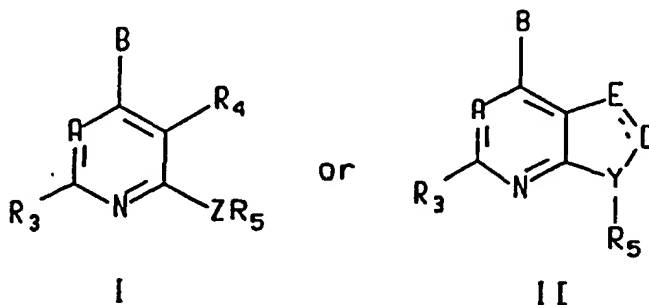
n-butyl-ethyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 di-n-propyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 ethyl-n-propyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 diethyl-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 5 n-butyl-ethyl-[2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 2-[N-n-butyl-N-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amino]-ethanol;
 4-(1-ethyl-propyl)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 n-butyl-ethyl-[2,5-dimethyl-7-(2,4-dimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-(1-ethyl-propyl)amine;
 10 2-[7-(4-bromo-2,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-(S)-[7-(4-bromo-2,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 4-(1-ethyl-propoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 4-(1-methoxymethyl-propoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 4-(1-ethyl-butyl)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo-[2,3-d]pyrimidine;
 15 [7-(4-bromo-2,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-(1-methoxymethyl-propyl)-
 amine;
 2-[7-(2-bromo-4,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(4-ethyl-2,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(2-ethyl-4,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 20 2-[7-(2-fluoromethyl-4,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol; and
 the pharmaceutically acceptable salts thereof.

20. A method of claim 2 wherein said corticotropin releasing factor antagonist is selected from the group consisting of:

4-(1-ethyl-propoxy)-2,5-dimethyl-6-(2,4,6-trimethyl-benzyl)-pyrimidine;
 2-(4-bromo-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 2-(4-ethyl-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 3-ethyl-4-(1-ethyl-propoxy)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
 2-(2,6-dimethyl-4-propyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 30 4-(1-ethyl-propoxy)-2-(4-methoxy-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridine;
 2-(4-ethoxy-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 2-(4-chloro-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
 4-(1-methoxymethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-diethyl-amine;
 35 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-propyl-amine;
 [2,5-dimethyl-6-(2,4,6-trimethyl-phenoxy)-pyrimidin-4-yl]-(1-ethyl-propyl)-amine;
 butyl-[3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-amine;
 4-(1-ethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethyl-phenylsulfanyl)-pyridine;
 butyl-[2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-ethyl-amine;
 40 4-(1-ethyl-propylamino)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-nicotinic acid methyl ester;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenylsulfanyl)-pyridin-4-yl]-ethyl-propyl-amine;
 [4-(1-ethyl-propylamino)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-3-yl]-methanol;
 [2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-ethyl-propyl-amine;
 1-(ethyl-propyl)-[6-methyl-3-nitro-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-amine;
 45 N4-(1-ethyl-propyl)-6-methyl-3-nitro-N2-(2,4,6-trimethyl-phenyl)-pyridine-2,4-diamine;
 N4-(1-ethyl-propyl)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
 N4-(1-ethyl-propyl)-6-methyl-N2-(2,4,6-trimethyl-phenyl)-pyridine-2,3,4-triamine;
 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-ethyl-(2,2,2-trifluoro-ethyl)-amine;
 [3-chloromethyl-6-methyl-2-(2,4,6-trimethyl-phenoxy)pyridin-4-yl]-(1-ethyl-propyl)-amine;
 50 [3,6-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-(1-ethyl-propyl)-amine;
 (1-ethyl-propyl)-[2-methyl-5-nitro-6-(2,4,6-trimethyl-pyridin-3-yloxy)-pyrimidin-4-yl]-amine;
 (1-ethyl-propyl)-[3-methoxymethyl-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridin-4-yl]-amine;
 N-(1-ethyl-propyl)-2-methyl-5-nitro-N-(2,4,6-trimethyl-pyridin-3-yl)-pyrimidine-4,6-diamine;
 [2-(4-chloro-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridin-4-yl]-diethyl-amine;
 55 4-(1-ethyl-propoxy)-3,6-dimethyl-2-(2,4,6-trimethylphenoxy)-pyridine;
 butyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-6,7-dihydro-5H-pyrrolo[2,3-d]pyrimidin-4-yl]-ethyl-amine;
 4-(butyl-ethylamino)-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-5,7-dihydro-pyrrolo[2,3-d]pyrimidin-6-one;
 4-(1-ethylpropoxy)-2,5-dimethyl-6-(2,4,6-trimethylphenoxy)-pyrimidine;

N-butyl-*N*-ethyl-2,5-dimethyl-*N*-(2,4,6-trimethylphenyl)-pyrimidine-4,6-diamine;
 (1-ethyl-propyl)-[5-methyl-3-(2,4,6-trimethyl-phenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-amine;
 [2,5-dimethyl-3-(2,4,6-trimethyl-phenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-(1-ethyl-propyl)-amine;
*N*4-(1-ethyl-propyl)-6, *N*3-dimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
 5 *N*4-(1-ethyl-propyl)-6, *N*3, *N*3-trimethyl-2-(2,4,6-trimethyl-phenoxy)-pyridine-3,4-diamine;
 6-(1-ethyl-propoxy)-2-methyl-*N*4-(2,4,6-trimethyl-phenyl)-pyrimidine-4,5-diamine;
 [4-(1-ethyl-propoxy)-3,6-dimethyl-pyridin-2-yl]-(2,4,6-trimethylphenyl)-amine;
 6-(ethyl-propyl-amino)-2,7-dimethyl-9-(2,4,6-trimethylphenyl)-7,9-dihydro-purin-8-one;
 butyl-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-ethylamine;
 10 3,6-dimethyl-4-(tetrahydrofuran-3-yloxy)-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridine
 [3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-(1-methoxymethylpropyl)-amine;
 4-(1-methoxymethylpropoxy)-3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridine;
 (1-ethylpropyl)-[3,5,6-trimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-b]pyridin-4-yl]-amine;
 4-(1-ethylpropoxy)-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-b]pyridine;
 15 4-(1-ethylpropoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-b]pyridine;
 4-(1-ethylpropoxy)-2,5-dimethyl-7-(2,6-dimethyl-4-bromophenyl)-7H-pyrrolo[2,3-b]pyridine;
 3-[(4-methyl-benzyl)-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amino]-propan-
 1-ol;
 diethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 20 2-butyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amino]-ethanol;
 dibutyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methyl-3-methylsulfonyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-cyclopropylmethyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-
 25 yl]-amine;
 di-1-propyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 diallyl-[6-methyl-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-chloro-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 butyl-ethyl-[6-methoxy-3-methylsulfanyl-1-(2,4,6-trichlorophenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 30 propyl-ethyl-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-amine;
 4-(1-ethyl-propyl)-6-methyl-3-methylsulfanyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidine;
 2-[3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamine]-butan-1-ol;
 [3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]-(1-methylpropyl)amine;
 4-(1-methoxymethylpropoxy)-3,6-dimethyl-1-(2,4,6-trimethylphenyl)-1H-pyrazolo[3,4-d]pyrimidine;
 35 *n*-butyl-ethyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 di-*n*-propyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 ethyl-*n*-propyl-[2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 diethyl-2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
n-butyl-ethyl-[2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 40 2-[*N*-*n*-butyl-*N*-(2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino]-ethanol;
 4-(1-ethyl-propyl)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
n-butyl-ethyl-[2,5-dimethyl-7-(2,4-dimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]amine;
 2,5-dimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-(1-ethyl-propyl)amine;
 2-[7-(4-bromo-2,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 45 2-(*S*)-[7-(4-bromo-2,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo(2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 4-(1-ethyl-propoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 4-(1-methoxymethyl-propoxy)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 4-(1-ethyl-butyl)-2,5,6-trimethyl-7-(2,4,6-trimethylphenyl)-7H-pyrrolo[2,3-d]pyrimidine;
 [7-(4-bromo-2,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-(1-methoxymethyl-propyl)-
 50 amine;
 2-[7-(2-bromo-4,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(4-ethyl-2,6-dimethyl-phenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(2-ethyl-4,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol;
 2-[7-(2-fluoromethyl-4,6-dimethylphenyl)-2,5-dimethyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino]-butan-1-ol and
 55 the pharmaceutically acceptable salts thereof;
 and said renin-angiotensin system inhibitor is selected from the group consisting of benazeprilat, captopril,
 enalapril, lisinopril, losartan, valsartan, irbesartan, candesartan, telmisartan, tasosartan and eprosartan.

21. A pharmaceutical composition which comprises amounts of a corticotropin releasing factor antagonist, a renin-angiotensin system inhibitor and a pharmaceutically acceptable carrier, vehicle or diluent.
22. A pharmaceutical composition which comprises congestive heart failure-treating or hypertension-treating amounts of a corticotropin releasing factor antagonist, a renin-angiotensin system inhibitor and a pharmaceutically acceptable carrier, vehicle or diluent.
23. A pharmaceutical composition of claim 22 wherein said corticotropin releasing factor antagonist and said renin-angiotensin system inhibitor are present in synergistic therapeutically effective amounts.
24. A pharmaceutical composition of claim 21 where said corticotropin releasing factor is a compound of structural formula



or a pharmaceutically acceptable salt thereof, wherein the dashed line represents an optional double bond:

A is $-CR_7$ or N;

B is $-\text{NR}_1\text{R}_2$, $-\text{CR}_1\text{R}_2\text{R}_{11}$, $-\text{C}(\text{=CR}_1\text{R}_2)\text{R}_2$, $-\text{NHCR}_{11}\text{R}_1\text{R}_2$, $-\text{OCR}_{11}\text{R}_1\text{R}_2$, $-\text{SCR}_{11}\text{R}_1\text{R}_2$, $-\text{CR}_{11}\text{R}_2\text{OR}_1$, $-\text{CR}_{11}\text{R}_2\text{SR}_1$, $-\text{C}(\text{S})\text{R}_2$, $-\text{NHNRR}_2$, $-\text{CR}_2\text{R}_{11}\text{NHR}_1$ or $-\text{C}(\text{O})\text{R}_2$;

D is: (i) N or -CR₁₀ when a double bond connects E and D and E is -CR₄; (ii) -CR₁₀ when a double bond connects E and D and E is N; or (iii) -CR₈R₉, -CHR₁₀, -C=O, -C=S, -C=NH, or -C=NCH₃ when a single bond connects E and D;

E is $-\text{CR}_4$ or N when a double bond connects E and D, and E is $-\text{CR}_4\text{R}_6$ or $-\text{NR}_6$ when a single bond connects E and D:

Y is N or-CH;

Z is NH, O, S, -N(C₁-C₂ alkyl) or -CR₁₂R₁₃, wherein R₁₂ and R₁₃ are each, independently, hydrogen, trifluoromethyl or methyl, or one of R₁₂ and R₁₃ is cyano and the other is hydrogen or methyl;

R₁ is hydrogen or C₁-C₆ alkyl which is optionally substituted with one or two substituents independently selected from hydroxy, cyano, nitro, fluoro, chloro, bromo, iodo, CF₃, C₁-C₄ alkoxy, -O-CO-(C₁-C₄ alkyl), -O-CO-NH(C₁-C₄ alkyl), -O-CO-N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -NH(C₁-C₄ alkyl), -N(C₁-C₂ alkyl)(C₁-C₄ alkyl), -S(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)CO(C₁-C₄ alkyl), -NHCO(C₁-C₄ alkyl), -CO₂(C₁-C₄ alkyl), -CONH(C₁-C₄ alkyl), -CON(C₁-C₄ alkyl)(C₁-C₂ alkyl), (C₁-C₄ alkyl)sulfinyl, (C₁-C₄ alkyl)sulfonyl, and (C₁-C₄ alkyl)sulfanyl, and wherein said C₁-C₆ alkyl, C₁-C₄ alkoxy and the C₁-C₄ alkyl moieties in the foregoing R₁ groups optionally contain one double or triple bond;

R₂ is C₁-C₆ alkyl, heteroaryl, aryl (heteroaryl)C₁-C₄ alkyl or (aryl)C₁-C₄ alkyl wherein said aryl and the aryl moiety of said (aryl)C₁-C₄ alkyl are selected from the group consisting of phenyl and naphthyl, and said heteroaryl and the heteroaryl moiety of said (heteroaryl)C₁-C₄ alkyl is selected from the group consisting of thienyl, benzothienyl, pyridyl, thiazolyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, furanyl, benzofuranyl, benzothiazolyl, isothiazolyl, benzisothiazolyl, benzisoxazolyl, benzimidazolyl, indolyl, and benzoxazolyl; or R² is C₃-C₈ cycloalkyl or (C₃-C₈ cycloalkyl)C₁-C₆ alkyl, wherein one or two of the ring carbons of said cycloalkyl having at least 4 ring members and the cycloalkyl moiety of said (C₃-C₈ cycloalkyl)C₁-C₆ alkyl having at least 4 ring members is optionally replaced with an oxygen or sulfur atom or with -NR₁₄ wherein R₁₄ is hydrogen or C₁-C₄ alkyl; and wherein each of the foregoing R₂ groups is optionally substituted with from one to three substituents independently selected from chloro, fluoro and C₁-C₄ alkyl, or with one substituent selected from bromo, iodo,

cyano, nitro, C₁-C₆ alkoxy, -O-CO-(C₁-C₄ alkyl), -O-CO-N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), (C₁-C₄ alkyl)sulfanyl, (C₁-C₄ alkyl)sulfinyl, and (C₁-C₄ alkyl)sulfonyl, and wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl and C₁-C₆ alkyl moieties of the foregoing R₂ groups optionally contain one carbon-carbon double or triple bond;

5 or R¹ and R² of said -NR₁R₂ and said -CR₁R₂R₁₁ are taken together to form a saturated 5 to 8 member ring, wherein said ring optionally contains one or two carbon-carbon double bonds, and wherein one or two of the ring carbons is optionally replaced with a heteroatom selected from O, S and N;

R₃ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, hydroxy, amino, SH, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CH₂OH, -CH₂OCH₃, -O(C₁-C₄ alkyl), (C₁-C₄ alkyl)sulfanyl, (C₁-C₄ alkyl)sulfonyl, or (C₁-C₄ alkyl)sulfinyl, wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl moieties of the foregoing R₃ groups optionally contain one double or triple bond and are optionally substituted with from one to three substituents independently selected from hydroxy, amino, C₁-C₃ alkoxy,

-NH(C₁-C₂ alkyl), -N(C₁-C₂)₂, -NHCOCH₃, fluoro, chloro and C₁-C₃ thioalkyl;

15 R₄ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, C₁-C₆ alkoxy, formyl, trifluoromethoxy, -CH₂OCH₃, -CH₂OCH₂CH₃, -CH₂CH₂OCH₃, -CH₂CF₃, CF₃, amino, nitro, -NH(C₁-C₄ alkyl), -N(CH₃)₂, -NHCOCH₃, -NHCONHCH₃, (C₁-C₄ alkyl)sulfanyl, (C₁-C₄ alkyl)sulfinyl, (C₁-C₄ alkyl)sulfonyl, cyano, hydroxy, -CO(C₁-C₄ alkyl), -CHO, or -CO₂(C₁-C₄ alkyl), wherein said C₁-C₆ alkyl, C₁-C₆ alkoxy and the C₁-C₄ alkyl moieties of the foregoing R₄ groups optionally contain one double or triple bond and are optionally substituted with one substituent selected from hydroxy, amino, -NHCOCH₃, -NH(C₁-C₂ alkyl), -N(C₁-C₂ alkyl)₂, -CO₂(C₁-C₄ alkyl), -CO(C₁-C₄ alkyl), C₁-C₃ alkoxy, (C₁-C₃ alkyl)sulfanyl, fluoro, chloro, cyano and nitro;

20 R₅ is phenyl, naphthyl, thienyl, benzothienyl, pyridyl, quinolyl, pyrazinyl, pyrimidyl, imidazolyl, furanyl, benzofuranyl, benzothiazolyl, isothiazolyl, benzoisothiazolyl, thiazolyl, isoxazolyl, benzisoxazolyl, benzimidazolyl, triazolyl, pyrazolyl, pyrrolyl, indolyl, azaindolyl, benzoxazolyl, oxazolyl, pyrrolidinyl, thiazolidinyl, morpholinyl, pyridinyl, tetrazolyl, or 3- to 8-membered cycloalkyl or 9- to 12-membered bicycloalkyl, wherein said cycloalkyl and bicycloalkyl optionally contain one or two of O, S or -N-G wherein G is hydrogen, C₁-C₄ alkyl, C₁-C₄ alkanoyl, phenyl or benzyl, wherein each of the above R₅ groups is optionally substituted with from one to three substituents independently selected from fluoro, chloro, C₁-C₆ alkyl, C₁-C₆ alkoxy and trifluoromethyl, or one substituent selected from bromo, iodo, cyano, nitro, amino, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), -CO(C₁-C₄ alkyl), -SO₂NH(C₁-C₄ alkyl), -SO₂N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -SO₂NH₂, -NHSO₂(C₁-C₄ alkyl), -S(C₁-C₄ alkyl), and -SO₂(C₁-C₄ alkyl), wherein said C₁-C₄ alkyl and C₁-C₆ alkyl moieties of the foregoing R₅ groups optionally contain one double or triple bond and are optionally substituted with one or two substituents independently selected from fluoro, chloro, hydroxy, amino, methylamino, dimethylamino and acetyl;

35 R₆ is hydrogen or C₁-C₆ alkyl, wherein said C₁-C₆ alkyl is optionally substituted with a single hydroxy, methoxy, ethoxy or fluoro group;

R₇ is hydrogen, C₁-C₄ alkyl, fluoro, chloro, bromo, iodo, cyano, hydroxy, C₁-C₄ alkoxy, -CO(C₁-C₄ alkyl), -CO₂(C₁-C₄ alkyl), -OCF₃, CF₃, -CH₂OH, -CH₂OCH₃ or -CH₂OCH₂CH₃;

R₈ and R₉ are each, independently, hydrogen, hydroxy, methyl, ethyl, methoxy, or ethoxy; or R₈ and R₉ together form an oxo (=O) group;

40 R₁₀ is hydrogen, C₁-C₆ alkyl, fluoro, chloro, bromo, iodo, C₁-C₆ alkoxy, formyl, amino, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -SO_n(C₁-C₄ alkyl), wherein n is 0, 1 or 2, cyano, carboxy, or amido, wherein said C₁-C₆ alkyl and the C₁-C₄ alkyl moieties of the foregoing R₁₀ groups are optionally substituted with one of hydroxy, trifluoromethyl, amino, carboxy, amido, -NHCO(C₁-C₄ alkyl), -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)(C₁-C₂ alkyl), -CO₂(C₁-C₄ alkyl), C₁-C₃ alkoxy, C₁-C₃ thioalkyl, fluoro, bromo, chloro, iodo, cyano or nitro; and

45 R₁₁ is hydrogen, hydroxy, fluoro, or methoxy; and

said renin-angiotensin system inhibitor is selected from the group consisting of a renin inhibitor, an angiotensin-converting enzyme inhibitor and an angiotensin II antagonist.

25. A composition of claim 24 wherein said compound of formula I or II is selected from the group consisting of:

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4(1-ethyl-propoxy)-2,5-dimethyl-6-(2,4,6-trimethyl-benzyl)-pyrimidine;
2-(4-bromo-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
2-(4-ethyl-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
3-ethyl-4-(1-ethyl-propoxy)-6-methyl-2-(2,4,6-trimethyl-phenoxy)-pyridine;
55 2-(2,6-dimethyl-4-propyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
4-(1-ethyl-propoxy)-2-(4-methoxy-2,6-dimethyl-phenoxy)-3,6-dimethyl-pyridine;
2-(4-ethoxy-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;
2-(4-chloro-2,6-dimethyl-phenoxy)-4-(1-ethyl-propoxy)-3,6-dimethyl-pyridine;